paragraphs (2) and (3) and inserting the following:

''(2) SPECIFIED TAX CREDIT BOND.—For purposes of this subsection, the term 'specified tax credit bond' means any qualified tax credit bond (as defined in section 54A(d)(1)) if the issuer of such bond makes an irrevocable election to have this subsection apply.''.

(B) Subparagraph (A) of section 6211(b)(4) of the Internal Revenue Code of 1986 is amended by striking ''and 6428A'' and inserting ''6428A, and 6431''.

SEC. 80704. GREEN BUILDING PRACTICES.

(a) IN GENERAL.—In carrying out a new construction or renovation project using any available project proceeds from the issuance of any qualified community college bond (as defined in subsection (a) of section 54 of the Internal Revenue Code of 1986), a governing body (as defined in subsection (c)(1) of such section) shall use, of those proceeds, not less than the applicable percentage described in subsection (b) for construction or renovation that is certified, verified, or consistent with the applicable provisions of—

(1) the Leadership in Energy and Environmental Design green building rating standard of the United States Green Building Council;

(2) the Living Building Challenge green building certification program developed by the International Living Future Institute;

(3) a green building rating program developed by the Collaborative for High-Performance Schools that is designated as CHPS Verified; or

(4) a green building program that—

(A) has standards that are equivalent to or more stringent than the standards of a program described in paragraph (1), (2), or (3);

(B) is adopted by the State or another jurisdiction with authority over the local educational agency; and

(C) includes a verifiable method to demonstrate compliance with the program.

(b) APPLICABLE PERCENTAGE DESCRIBED.—The applicable percentage referred to in subsection (a) is—

(1) for fiscal year 2022, 60 percent;

(2) for fiscal year 2023, 70 percent;

(3) for fiscal year 2024, 80 percent;

(4) for fiscal year 2025, 90 percent; and

(5) for each of fiscal years 2026 through 2031, 100 percent.

SEC. 80705. USE OF AMERICAN IRON, STEEL, AND MANUFACTURED PRODUCTS.

(a) IN GENERAL.—A governing body (as defined in subsection (c)(1) of section 54 of the Internal Revenue Code of 1986) that receives covered funds shall ensure that any iron, steel, and manufactured products used in projects carried out with such funds are produced in the United States.

(b) WAIVER AUTHORITY.—

(1) IN GENERAL.—The Secretary may waive the requirement of subsection (a) if the Secretary determines that—

(A) applying subsection (a) would be inconsistent with the public interest;

(B) iron, steel, and manufactured products produced in the United States are not produced in a sufficient and reasonably available amount or are not of a satisfactory quality; or

(C) using iron, steel, and manufactured products produced in the United States will increase the cost of the overall project by more than 25 percent.

(2) PUBLICATION.—Before issuing a waiver under paragraph (1), the Secretary shall publish in the Federal Register a detailed written explanation of the waiver determination.

(c) CONSISTENCY WITH INTERNATIONAL AGREEMENTS.—This section shall be applied in a manner consistent with the obligations of the United States under international agreements.

(d) DEFINITIONS.—In this section:

(1) COVERED FUNDS.—The term ''covered funds'' means any available project proceeds from the issuance of any qualified community college bond (as defined in section 54(a) of the Internal Revenue Code of 1986).

(2) MANUFACTURED PRODUCT.—The term ''manufactured product'' means any construction material or end product (as such terms are defined in part 25.003 of the Federal Acquisition Regulation) that is not an iron or steel product, including—

(A) electrical components; and

(B) non-ferrous building materials, including aluminum and polyvinylchloride (PVC), glass, fiber optics, plastic, wood, masonry, rubber, manufactured stone, any other nonferrous metals, and any unmanufactured construction material.

(3) PRODUCED IN THE UNITED STATES.—The term ''produced in the United States'' means the following:

(A) When used with respect to a manufactured product, the product was manufactured in the United States and the cost of the components of such product that were mined, produced, or manufactured in the United States exceeds 60 percent of the total cost of all components of the product.

(B) When used with respect to iron or steel products, or an individual component of a manufactured product, all manufacturing processes for such iron or steel products or components, from the initial melting stage through the application of coatings, occurred in the United States, except that the term does not include—

(i) steel or iron material or products manufactured abroad from semi-finished steel or iron from the United States; and

(ii) steel or iron material or products manufactured in the United States from semi-finished steel or iron of foreign origin.

(4) SECRETARY.—The term ''Secretary'' means the Secretary of Education.

SEC. 80706. EFFECTIVE DATE.

The amendments made by this title shall apply to obligations issued after the date of the enactment of this Act.

SA 2550. Mr. OSSOFF (for himself and Mr. WHITEHOUSE) submitted an amendment intended to be proposed to amendment SA 2137 proposed by Mr. SCHUMER (for Ms. SINEMA (for herself, Mr. PORTMAN, Mr. MANCHIN, Mr. CASSIDY, Mrs. SHAHEEN, Ms. COLLINS, Mr. TESTER, Ms. MURKOWSKI, Mr. WARNER, and Mr. ROMNEY)) to the bill H.R. 3684, to authorize funds for Federal-aid highways, highway safety programs, and transit programs, and for other purposes; which was ordered to lie on the table; as follows:

On page 202, strike line 5 and insert the following:

(2) in subsection (l)—

(A) by striking paragraph (2);

(B) by striking the subsection designation and all that follows through ''In determining'' in paragraph (1) in the matter preceding subparagraph (A) and inserting the following:

''(l) ACCOMMODATING UTILITY FACILITIES IN THE RIGHT-OF-WAY.—

''(1) DEFINITIONS.—In this subsection:

''(A) FEDERAL LAND.—The term 'Federal land' means any land or interest in land owned by the United States.

''(B) INDIAN LAND.—The term 'Indian land' means—

''(i) land within the limits of an Indian reservation; or

''(ii) land over which an Indian Tribe exercises governmental power and that is—

''(I) held in trust by the United States for the benefit of any Indian tribe or individual Indian; or

''(II) held by an Indian Tribe or individual Indian subject to restriction by the United States against alienation.

''(C) RIGHT-OF-WAY.—The term 'right-of-way' means any real property, or interest therein, acquired, dedicated, or reserved for the construction, operation, and maintenance of a highway.

''(D) UTILITY FACILITY.—

''(i) IN GENERAL.—The term 'utility facility' means any privately, publicly, or cooperatively owned line, facility, or system for producing, transmitting, or distributing communications, power, electricity, light, heat, gas, water, steam, waste, storm water not connected with highway drainage, or any other similar commodity, including any fire or police signal system or street lighting system, that directly or indirectly serves the public.

''(ii) INCLUSIONS.—The term 'utility facility' includes—

''(I) a renewable energy generation facility;

''(II) electrical transmission and distribution infrastructure; and

''(III) broadband infrastructure and conduit.

''(2) ACCOMMODATION.—In determining''; and

(C) by adding at the end the following:

''(3) STATE APPROVAL.—A State, on behalf of the Secretary, may approve accommodating a utility facility described in paragraph (1)(D)(ii) within a right-of-way on a Federal-aid highway.

''(4) EXCLUSION.—Paragraph (3) shall not apply to a utility facility on Federal land or Indian land.

''(5) SAVINGS PROVISION.—Nothing in this subsection alters or affects any prohibition relating to commercial activity under section 111(a).'';

(3) in subsection (o)—

On page 202, line 23, strike ''(3)'' and insert ''(4)''.

On page 203, strike line 17 and insert the following:

the project is located on a Federal-aid highway.

''(t) VEGETATION MANAGEMENT.—States are encouraged to implement, or to enter into partnerships to implement, vegetation management practices, such as increased mowing heights and planting native grasses and pollinator-friendly habitats, along a right-of-way on a Federal-aid highway, if the implementation of those practices—

''(1) is in the public interest; and

''(2) will not impair the highway or interfere with the free and safe flow of traffic.''.

SA 2551. Mrs. FEINSTEIN submitted an amendment intended to be proposed to amendment SA 2137 proposed by Mr. SCHUMER (for Ms. SINEMA (for herself, Mr. PORTMAN, Mr. MANCHIN, Mr. CASSIDY, Mrs. SHAHEEN, Ms. COLLINS, Mr. TESTER, Ms. MURKOWSKI, Mr. WARNER, and Mr. ROMNEY)) to the bill H.R. 3684, to authorize funds for Federal-aid highways, highway safety programs, and transit programs, and for other purposes; which was ordered to lie on the table; as follows:

On page 1200, strike line 9, and all that follows through page 1202, line 10, and insert the following:

Subtitle B—Cannabidiol and Marihuana Research Expansion

SEC. 25101. SHORT TITLE.

This subtitle may be cited as the ''Cannabidiol and Marihuana Research Expansion Act''.

SEC. 25102. DEFINITIONS.

In this subtitle—

(1) the term ''appropriately registered'' means that an individual or entity is registered under the Controlled Substances Act (21 U.S.C. 801 et seq.) to engage in the type of activity that is carried out by the individual or entity with respect to a controlled substance on the schedule that is applicable to cannabidiol or marihuana, as applicable;

(2) the term ''cannabidiol'' means—

(A) the substance, cannabidiol, as derived from marihuana that has a delta-9-tetrahydrocannabinol level that is greater than 0.3 percent; and

(B) the synthetic equivalent of the substance described in subparagraph (A);

(3) the terms ''controlled substance'', ''dispense'', ''distribute'', ''manufacture'', ''marihuana'', and ''practitioner'' have the meanings given such terms in section 102 of the Controlled Substances Act (21 U.S.C. 802), as amended by this subtitle;

(4) the term ''covered institution of higher education'' means an institution of higher education (as defined in section 101 of the Higher Education Act of 1965 (20 U.S.C. 1001)) that—

(A)(i) has highest or higher research activity, as defined by the Carnegie Classification of Institutions of Higher Education; or

(ii) is an accredited medical school or an accredited school of osteopathic medicine; and

(B) is appropriately registered under the Controlled Substances Act (21 U.S.C. 801 et seq.);

(5) the term ''drug'' has the meaning given the term in section 201(g)(1) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321(g)(1));

(6) the term ''medical research for drug development'' means medical research that is—

(A) a preclinical study or clinical investigation conducted in accordance with section 505(i) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(i)) or otherwise permitted by the Department of Health and Human Services to determine the potential medical benefits of marihuana or cannabidiol as a drug; and

(B) conducted by a covered institution of higher education, practitioner, or manufacturer that is appropriately registered under the Controlled Substances Act (21 U.S.C. 801 et seq.); and

(7) the term ''State'' means any State of the United States, the District of Columbia, and any territory of the United States.

CHAPTER 1—REGISTRATIONS FOR MARIHUANA RESEARCH

SEC. 25121. MARIHUANA RESEARCH APPLICATIONS.

Section 303(f) of the Controlled Substances Act (21 U.S.C. 823(f)) is amended—

(1) by redesignating paragraphs (1) through (5) as subparagraphs (A) through (E), respectively;

(2) by striking ''(f) The Attorney General'' and inserting ''(f)(1) The Attorney General'';

(3) by striking ''Registration applications'' and inserting the following:

''(2)(A) Registration applications'';

(4) by striking ''Article 7'' and inserting the following:

''(3) Article 7''; and

(5) by inserting after paragraph (2)(A), as so designated, the following:

''(B)(i) The Attorney General shall register a practitioner to conduct research with marihuana if—

''(I) the applicant's research protocol—

''(aa) has been reviewed and allowed—

''(AA) by the Secretary of Health and Human Services under section 505(i) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(i));

''(BB) by the National Institutes of Health or another Federal agency that funds scientific research; or

''(CC) pursuant to sections 1301.18 and 1301.32 of title 21, Code of Federal Regulations, or any successors thereto; and

''(II) the applicant has demonstrated to the Attorney General that there are effective procedures in place to adequately safeguard against diversion of the controlled substance for legitimate medical or scientific use pursuant to section 25125 of the Cannabidiol and Marihuana Research Expansion Act, including demonstrating that the security measures are adequate for storing the quantity of marihuana the applicant would be authorized to possess.

''(ii) The Attorney General may deny an application for registration under this subparagraph only if the Attorney General determines that the issuance of the registration would be inconsistent with the public interest. In determining the public interest, the Attorney General shall consider the factors listed in—

''(I) subparagraphs (B) through (E) of paragraph (1); and

''(II) subparagraph (A) of paragraph (1), if the applicable State requires practitioners conducting research to register with a board or authority described in such subparagraph (A).

''(iii)(I) Not later than 60 days after the date on which the Attorney General receives a complete application for registration under this subparagraph, the Attorney General shall—

''(aa) approve the application; or

''(bb) request supplemental information.

''(II) For purposes of subclause (I), an application shall be deemed complete when the applicant has submitted documentation showing that the requirements under clause (i) are satisfied.

''(iv) Not later than 30 days after the date on which the Attorney General receives supplemental information as described in clause (iii)(I)(bb) in connection with an application described in this subparagraph, the Attorney General shall approve or deny the application.

''(v) If an application described in this subparagraph is denied, the Attorney General shall provide a written explanation of the basis of denial to the applicant.''.

SEC. 25122. RESEARCH PROTOCOLS.

(a) IN GENERAL.—Paragraph (2)(B) of section 303(f) of the Controlled Substances Act (21 U.S.C. 823(f)), as amended by section 25121 of this Act, is further amended by adding at the end the following:

''(vi)(I) If the Attorney General grants an application for registration under clause (i), the registrant may amend or supplement the research protocol without reapplying if the registrant does not change—

''(aa) the quantity or type of drug;

''(bb) the source of the drug; or

''(cc) the conditions under which the drug is stored, tracked, or administered.

''(II)(aa) If a registrant under clause (i) seeks to change the type of drug, the source of the drug, or conditions under which the drug is stored, tracked, or administered, the registrant shall notify the Attorney General via registered mail, or an electronic means permitted by the Attorney General, not later than 30 days before implementing an amended or supplemental research protocol.

''(bb) A registrant may proceed with an amended or supplemental research protocol described in item (aa) if the Attorney General does not explicitly object during the 30-day period beginning on the date on which the Attorney General receives the notice under item (aa).

''(cc) The Attorney General may only object to an amended or supplemental research protocol under this subclause if additional security measures are needed to safeguard against diversion or abuse.

''(dd) If a registrant under clause (i) seeks to address additional security measures identified by the Attorney General under item (cc), the registrant shall notify the Attorney General via registered mail, or an electronic means permitted by the Attorney General, not later than 30 days before implementing an amended or supplemental research protocol.

''(ee) A registrant may proceed with an amended or supplemental research protocol described in item (dd) if the Attorney General does not explicitly object during the 30-day period beginning on the date on which the Attorney General receives the notice under item (dd).

''(III)(aa) If a registrant under clause (i) seeks to change the quantity of marihuana needed for research and the change in quantity does not impact the factors described in item (bb) or (cc) of subclause (I) of this clause, the registrant shall notify the Attorney General via registered mail or using an electronic means permitted by the Attorney General.

''(bb) A notification under item (aa) shall include—

''(AA) the Drug Enforcement Administration registration number of the registrant;

''(BB) the quantity of marihuana already obtained;

''(CC) the quantity of additional marihuana needed to complete the research; and

''(DD) an attestation that the change in quantity does not impact the source of the drug or the conditions under which the drug is stored, tracked, or administered.

''(cc) The Attorney General shall ensure that—

''(AA) any registered mail return receipt with respect to a notification under item (aa) is submitted for delivery to the registrant providing the notification not later than 3 days after receipt of the notification by the Attorney General; and

''(BB) notice of receipt of a notification using an electronic means permitted under item (aa) is provided to the registrant providing the notification not later than 3 days after receipt of the notification by the Attorney General.

''(dd)(AA) On and after the date described in subitem (BB), a registrant that submits a notification in accordance with item (aa) may proceed with the research as if the change in quantity has been approved on such date, unless the Attorney General notifies the registrant of an objection described in item (ee).

''(BB) The date described in this subitem is the date on which a registrant submitting a notification under item (aa) receives the registered mail return receipt with respect to the notification or the date on which the registrant receives notice that the notification using an electronic means permitted under item (aa) was received by the Attorney General, as the case may be.

''(ee) A notification submitted under item (aa) shall be deemed to be approved unless the Attorney General, not later than 10 days after receiving the notification, explicitly objects based on a finding that the change in quantity—

''(AA) does impact the source of the drug or the conditions under which the drug is stored, tracked, or administered; or

''(BB) necessitates that the registrant implement additional security measures to safeguard against diversion or abuse.

''(IV) Nothing in this clause shall limit the authority of the Secretary of Health and Human Services over requirements related to research protocols, including changes in—

''(aa) the method of administration of marihuana;

''(bb) the dosing of marihuana; and

''(cc) the number of individuals or patients involved in research.''.

(b) REGULATIONS.—Not later than 1 year after the date of enactment of this Act, the Attorney General shall promulgate regulations to carry out the amendment made by this section.

SEC. 25123. APPLICATIONS TO MANUFACTURE MARIHUANA FOR RESEARCH.

(a) IN GENERAL.—Section 303 of the Controlled Substances Act (21 U.S.C. 823) is amended—

(1) by redesignating subsections (c) through (k) as subsections (d) through (l), respectively;

(2) by inserting after subsection (b) the following:

''(c)(1)(A) As it relates to applications to manufacture marihuana for research purposes, if the Attorney General places a notice in the Federal Register to increase the number of entities registered under this Act to manufacture marihuana to supply appropriately registered researchers in the United States, the Attorney General shall, not later than 60 days after the date on which the Attorney General receives a completed application—

''(i) approve the application; or

''(ii) request supplemental information.

''(B) For purposes of subparagraph (A), an application shall be deemed complete when the applicant has submitted documentation showing each of the following:

''(i) The requirements designated in the notice in the Federal Register are satisfied.

''(ii) The requirements under this Act are satisfied.

''(iii) The applicant will limit the transfer and sale of any marihuana manufactured under this subsection—

''(I) to researchers who are registered under this Act to conduct research with controlled substances in schedule I; and

''(II) for purposes of use in preclinical research or in a clinical investigation pursuant to an investigational new drug exemption under 505(i) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(i)).

''(iv) The applicant will transfer or sell any marihuana manufactured under this subsection only with prior, written consent for the transfer or sale by the Attorney General.

''(v) The applicant has completed the application and review process under subsection (a) for the bulk manufacture of controlled substances in schedule I.

''(vi) The applicant has established and begun operation of a process for storage and handling of controlled substances in schedule I, including for inventory control and monitoring security in accordance with section 25125 of the Cannabidiol and Marihuana Research Expansion Act.

''(vii) The applicant is licensed by each State in which the applicant will conduct operations under this subsection, to manufacture marihuana, if that State requires such a license.

''(C) Not later than 30 days after the date on which the Attorney General receives supplemental information requested under subparagraph (A)(ii) with respect to an application, the Attorney General shall approve or deny the application.

''(2) If an application described in this subsection is denied, the Attorney General shall provide a written explanation of the basis of denial to the applicant.'';

(3) in subsection (h)(2), as so redesignated, by striking ''subsection (f)'' each place it appears and inserting ''subsection (g)'';

(4) in subsection (j)(1), as so redesignated, by striking ''subsection (d)'' and inserting ''subsection (e)''; and

(5) in subsection (k), as so redesignated, by striking ''subsection (f)'' each place it appears and inserting ''subsection (g)''.

(b) TECHNICAL AND CONFORMING AMENDMENTS.—

(1) The Controlled Substances Act (21 U.S.C. 801 et seq.) is amended—

(A) in section 102 (21 U.S.C. 802)—

(i) in paragraph (16)(B)—

(I) in clause (i), by striking ''or'' at the end;

(II) by redesignating clause (ii) as (iii); and

(III) by inserting after clause (i) the following:

''(ii) the synthetic equivalent of hemp-derived cannabidiol that contains less than 0.3 percent tetrahydrocannabinol; or'';

(ii) in paragraph (52)(B)—

(I) by striking ''303(f)'' each place it appears and inserting ''303(g)''; and

(II) in clause (i), by striking ''(d), or (e)'' and inserting ''(e), or (f)''; and

(iii) in paragraph (54), by striking ''303(f)'' each place it appears and inserting ''303(g)'';

(B) in section 302(g)(5)(A)(iii)(I)(bb) (21 U.S.C. 822(g)(5)(A)(iii)(I)(bb)), by striking ''303(f)'' and inserting ''303(g)'';

(C) in section 304 (21 U.S.C. 824), by striking ''303(g)(1)'' each place it appears and inserting ''303(h)(1)'';

(D) in section 307(d)(2) (21 U.S.C. 827(d)(2)), by striking ''303(f)'' and inserting ''303(g)'';

(E) in section 309A(a)(2) (21 U.S.C. 829a(a)(2)), in the matter preceding subparagraph (A), by striking ''303(g)(2)'' and inserting ''303(h)(2)'';

(F) in section 311(h) (21 U.S.C. 831(h)), by striking ''303(f)'' each place it appears and inserting ''303(g)'';

(G) in section 401(h)(2) (21 U.S.C. 841(h)(2)), by striking ''303(f)'' each place it appears and inserting ''303(g)'';

(H) in section 403(c)(2)(B) (21 U.S.C. 843(c)(2)(B)), by striking ''303(f)'' and inserting ''303(g)''; and

(I) in section 512(c)(1) (21 U.S.C. 882(c)(1)) by striking ''303(f)'' and inserting ''303(g)''.

(2) Section 1008(c) of the Controlled Substances Import and Export Act (21 U.S.C. 958(c)) is amended—

(A) in paragraph (1), by striking ''303(d)'' and inserting ''303(e)''; and

(B) in paragraph (2)(B), by striking ''303(h)'' and inserting ''303(i)''.

(3) Title V of the Public Health Service Act (42 U.S.C. 290aa et seq.) is amended—

(A) in section 520E–4(c) (42 U.S.C. 290bb–36d(c)), by striking ''303(g)(2)(B)'' and inserting ''303(h)(2)(B)''; and

(B) in section 544(a)(3) (42 U.S.C. 290dd–3(a)(3)), by striking ''303(g)'' and inserting ''303(h)''.

(4) Title XVIII of the Social Security Act (42 U.S.C. 1395 et seq.) is amended—

(A) in section 1833(bb)(3)(B) (42 U.S.C. 1395l(bb)(3)(B)), by striking ''303(g)'' and inserting ''303(h)'';

(B) in section 1834(o)(3)(C)(ii) (42 U.S.C. 1395m(o)(3)(C)(ii)), by striking ''303(g)'' and inserting ''303(h)''; and

(C) in section 1866F(c)(3)(C) (42 U.S.C. 1395cc–6(c)(3)(C)), by striking ''303(g)'' and inserting ''303(h)''.

(5) Section 1903(aa)(2)(C)(ii) of the Social Security Act (42 U.S.C. 1396b(aa)(2)(C)(ii)) is amended by striking ''303(g)'' each place it appears and inserting ''303(h)''.

SEC. 25124. ADEQUATE AND UNINTERRUPTED SUPPLY.

On an annual basis, the Attorney General shall assess whether there is an adequate and uninterrupted supply of marihuana, including of specific strains, for research purposes.

SEC. 25125. SECURITY REQUIREMENTS.

(a) IN GENERAL.—An individual or entity engaged in researching marihuana or its components shall store it in a securely locked, substantially constructed cabinet.

(b) REQUIREMENTS FOR OTHER MEASURES.—Any other security measures required by the Attorney General to safeguard against diversion shall be consistent with those required for practitioners conducting research on other controlled substances in schedules I and II in section 202(c) of the Controlled Substances Act (21 U.S.C. 812(c)) that have a similar risk of diversion and abuse.

SEC. 25126. PROHIBITION AGAINST REINSTATING INTERDISCIPLINARY REVIEW PROCESS FOR NON-NIH-FUNDED RESEARCHERS.

The Secretary of Health and Human Services may not—

(1) reinstate the Public Health Service interdisciplinary review process described in the guidance entitled ''Guidance on Procedures for the Provision of Marijuana for Medical Research'' (issued on May 21, 1999); or

(2) require another review of scientific protocols that is applicable only to research on marihuana or its components.

CHAPTER 2—DEVELOPMENT OF FDA-APPROVED DRUGS USING CANNABIDIOL AND MARIHUANA

SEC. 25141. MEDICAL RESEARCH ON CANNABIDIOL.

Notwithstanding any provision of the Controlled Substances Act (21 U.S.C. 801 et seq.), the Safe and Drug-Free Schools and Communities Act (20 U.S.C. 7101 et seq.), chapter 81 of title 41, United States Code, or any other Federal law, an appropriately registered covered institution of higher education, a practitioner, or a manufacturer may manufacture, distribute, dispense, or possess marihuana or cannabidiol if the marihuana or cannabidiol is manufactured, distributed, dispensed, or possessed, respectively, for purposes of medical research for drug development or subsequent commercial production in accordance with section 25142.

SEC. 25142. REGISTRATION FOR THE COMMERCIAL PRODUCTION AND DISTRIBUTION OF FOOD AND DRUG ADMINISTRATION-APPROVED DRUGS.

The Attorney General shall register an applicant to manufacture or distribute cannabidiol or marihuana for the purpose of commercial production of a drug containing or derived from marihuana that is approved by the Secretary of Health and Human Services under section 505 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355), in accordance with the applicable requirements under subsection (a) or (b) of section 303 of the Controlled Substances Act (21 U.S.C. 823).

SEC. 25143. IMPORTATION OF CANNABIDIOL FOR RESEARCH PURPOSES.

The Controlled Substances Import and Export Act (21 U.S.C. 951 et seq.) is amended—

(1) in section 1002(a) (21 U.S.C. 952(a))—

(A) in paragraph (1), by striking ''and'' at the end;

(B) in paragraph (2)(C), by inserting ''and'' after ''uses,''; and

(C) inserting before the undesignated matter following paragraph (2)(C) the following:

''(3) such amounts of marihuana or cannabidiol (as defined in section 25102 of the Cannabidiol and Marihuana Research Expansion Act) as are—

''(A) approved for medical research for drug development (as such terms are defined in section 25102 of the Cannabidiol and Marihuana Research Expansion Act), or

''(B) necessary for registered manufacturers to manufacture drugs containing marihuana or cannabidiol that have been approved for use by the Commissioner of Food and Drugs under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.),''; and

(2) in section 1007 (21 U.S.C. 957), by amending subsection (a) to read as follows:

''(a)(1) Except as provided in paragraph (2), no person may—

''(A) import into the customs territory of the United States from any place outside thereof (but within the United States), or import into the United States from any place outside thereof, any controlled substance or list I chemical, or

''(B) export from the United States any controlled substance or list I chemical,

unless there is in effect with respect to such person a registration issued by the Attorney General under section 1008, or unless such person is exempt from registration under subsection (b).

''(2) Paragraph (1) shall not apply to the import or export of marihuana or cannabidiol (as defined in section 25102 of the Cannabidiol and Marihuana Research Expansion Act) that has been approved for—

''(A) medical research for drug development authorized under section 25141 of the Cannabidiol and Marihuana Research Expansion Act; or

''(B) use by registered manufacturers to manufacture drugs containing marihuana or cannabidiol that have been approved for use by the Commissioner of Food and Drugs under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.).''.

CHAPTER 3—DOCTOR-PATIENT RELATIONSHIP

SEC. 25161. DOCTOR-PATIENT RELATIONSHIP.

It shall not be a violation of the Controlled Substances Act (21 U.S.C. 801 et seq.) for a State-licensed physician to discuss—

(1) the currently known potential harms and benefits of marihuana derivatives, including cannabidiol, as a treatment with the legal guardian of the patient of the physician if the patient is a child; or

(2) the currently known potential harms and benefits of marihuana and marihuana derivatives, including cannabidiol, as a treatment with the patient or the legal guardian of the patient of the physician if the patient is a legal adult.

CHAPTER 4—FEDERAL RESEARCH

SEC. 25181. FEDERAL RESEARCH.

(a) IN GENERAL.—Not later than 1 year after the date of enactment of this Act, the Secretary of Health and Human Services, in coordination with the Director of the National Institutes of Health and the heads of other relevant Federal agencies, shall submit to the Caucus on International Narcotics Control, the Committee on the Judiciary, and the Committee on Health, Education, Labor, and Pensions of the Senate and the Committee on Energy and Commerce and the Committee on the Judiciary of the House of Representatives a report on—

(1) the potential therapeutic effects of cannabidiol or marihuana on serious medical conditions, including intractable epilepsy;

(2) the potential effects of marihuana, including—

(A) the effect of increasing delta-9-tetrahydrocannabinol levels on the human body and developing adolescent brains; and

(B) the effect of various delta-9-tetrahydrocannabinol levels on cognitive abilities, such as those that are required to operate motor vehicles or other heavy equipment; and

(3) the barriers associated with researching marihuana or cannabidiol in States that have legalized the use of such substances, which shall include—

(A) recommendations as to how such barriers might be overcome, including whether public-private partnerships or Federal-State research partnerships may or should be implemented to provide researchers with access to additional strains of marihuana and cannabidiol; and

(B) recommendations as to what safeguards must be in place to verify—

(i) the levels of tetrahydrocannabinol, cannabidiol, or other cannabinoids contained in products obtained from such States is accurate; and

(ii) that such products do not contain harmful or toxic components.

(b) ACTIVITIES.—To the extent practicable, the Secretary of Health and Human Services, either directly or through awarding grants, contacts, or cooperative agreements, shall expand and coordinate the activities of the National Institutes of Health and other relevant Federal agencies to better determine the effects of cannabidiol and marihuana, as outlined in the report submitted under paragraphs (1) and (2) of subsection (a).

Subtitle C—GAO Study

SEC. 25201. GAO STUDY ON IMPROVING THE EFFICIENCY OF TRAFFIC SYSTEMS.

Not later than 1 year after the date of enactment of this Act, the Comptroller General of the United States shall carry out, and submit to Congress a report describing the results of, a study on the potential societal benefits of improving the efficiency of traffic systems.

SA 2552. Mrs. MURRAY (for herself and Ms. CORTEZ MASTO) submitted an amendment intended to be proposed to amendment SA 2137 proposed by Mr. SCHUMER (for Ms. SINEMA (for herself, Mr. PORTMAN, Mr. MANCHIN, Mr. CASSIDY, Mrs. SHAHEEN, Ms. COLLINS, Mr. TESTER, Ms. MURKOWSKI, Mr. WARNER, and Mr. ROMNEY)) to the bill H.R. 3684, to authorize funds for Federal-aid highways, highway safety programs, and transit programs, and for other purposes; which was ordered to lie on the table; as follows:

On page 2386, line 17, strike ''or in part''.

SA 2553. Mr. HEINRICH (for himself, Mr. MORAN, and Mr. LUJÁN) submitted an amendment intended to be proposed to amendment SA 2137 proposed by Mr. SCHUMER (for Ms. SINEMA (for herself, Mr. PORTMAN, Mr. MANCHIN, Mr. CASSIDY, Mrs. SHAHEEN, Ms. COLLINS, Mr. TESTER, Ms. MURKOWSKI, Mr. WARNER, and Mr. ROMNEY)) to the bill H.R. 3684, to authorize funds for Federal-aid highways, highway safety programs, and transit programs, and for other purposes; which was ordered to lie on the table; as follows:

At the appropriate place in division G, insert the following:

TITLE ____—CHAMPIONING APPRENTICESHIPS FOR NEW CAREERS AND EMPLOYEES IN TECHNOLOGY

SEC. _____1. SHORT TITLE.

This title may be cited as the ''Championing Apprenticeships for New Careers and Employees in Technology Act'' or the ''CHANCE in TECH Act''.

SEC. _____2. CONGRESSIONAL FINDINGS.

Congress finds the following:

(1) During any given 90-day period there can be more than 500,000 information technology job openings in the United States.

(2) Employment in the technology sector is growing twice as fast as employment in the United States.

(3) Jobs in the technology sector tend to provide higher pay and better benefits than other jobs and have been more resilient to economic downturn than jobs available in other private sector industries.

(4) Information technology skills are transferrable across nearly all industries.

(5) Exceptional education and on-the-job training programs exist and should be scaled to meet the demands of the modern technology workforce.

(6) Adoption of existing employer-driven intermediary models, such as ApprenticeshipUSA under the Department of Labor, will help grow the information technology workforce.

(7) Career pathway education should start in high school through pathways and programs of study that align with local and regional employer needs.

(8) Preparing a student for a job in the technology sector is essential to the growth and competitiveness of the economy in the United States in the 21st Century.

(9) Nearly 800,000 information technology workers will retire between 2017 and 2024.

(10) According to the Bureau of Labor Statistics, in May 2020, the median annual wage for computer and information technology occupations was $91,250, which was higher than the median annual wage for all occupations of $41,950.

SEC. _____3. TECHNOLOGY APPRENTICESHIP CONTRACTS.

(a) IN GENERAL.—The Secretary of Labor (referred to in this section as ''the Secretary'') shall enter into contracts with industry intermediaries for the purpose of promoting the development of and access to apprenticeships in the technology sector, from amounts appropriated under subsection (e).

(b) ELIGIBILITY.—To be eligible to be awarded a contract under this section, an industry intermediary shall submit an application to the Secretary, at such time and in such a manner as may be required by the Secretary, that identifies proposed activities designed to further the purpose described in subsection (a).

(c) SELECTION.—The Secretary shall award contracts under this section based on competitive criteria to be prescribed by the Secretary.

(d) CONTRACTOR ACTIVITIES.—An industry intermediary that is awarded a contract under this section may only use the funds made available through such contract to carry out activities designed to further the purpose described in subsection (a), including—

(1) facilitating the provision and development of apprenticeships in the technology sector through collaborations with public and private entities that provide job-related instruction, such as on-the-job training, pre-apprenticeship training, and technical training;

(2) encouraging entities to establish such apprenticeships;

(3) identifying, assessing, and training applicants for such apprenticeships who are—

(A) enrolled in high school;

(B) enrolled in an early college high school that focuses on education in STEM subjects;

(C) individuals aged 18 years or older who meet appropriate qualification standards; or

(D) enrolled in pre-apprenticeship or apprenticeship training initiatives that allow adults to concurrently increase academic and workforce skills through proven, evidence-based models that connect all learning to the specific apprenticeship involved and significantly accelerate completion of preparation for the apprenticeship; and

(4) tracking the progress of such applicants who participate in such apprenticeships.

(e) AUTHORIZATION OF APPROPRIATIONS.—There are authorized to be appropriated to